



Plant

- Morphine, heroin,codeine—opium plant
- Atropine—Atropabelladonna
- Digoxin—purple foxglove





Animal

- Insulin/Oxytocin—beef/pork/human
- Cod-liver oil--Fish

Mineral

- Sodium bicarbonate
- Calcium Chloride
- Magnesium Sulfate



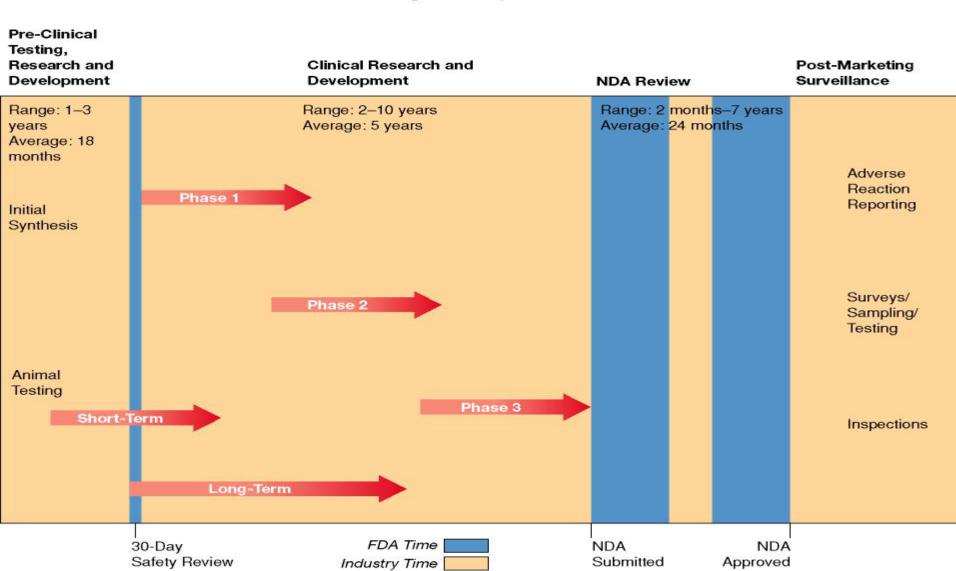


• Synthetic

- Diazepam
- Human Insulin

Bringing New Drugs to Market

New Drug Development Timeline



Phases of Human Studies

Phase 1

 To determine the drug's pharmacokinetics, toxicity, and safe dose in humans. Limited populations of healthy human volunteers

Phase 2

 To determine the therapeutic drug level and watch for toxic side effects. Tested on limited populations of patients who have the disease it is intended to treat.

Phases of Human Studies (cont)

Phase 3

To refine the usual therapeutic dose and collect data on side effects. Requires a larger patient population. Studies are usually double blind

Phase 4

Involves post-marketing analysis during conditional approval

FDA Classification of New Drugs

- Utilizes a number and a letter for each new drug
 - Numerical Classification (Chemical)
 - Assigned a number 1- 7
 - Letter Classification (Treatment or Therapeutic Potential)
- Other Classification

Laws Affecting Drug Administration

- 1906: Pure Food & Drug Act
 - Enacted to improve the quality and labeling of drugs
 - Established USP and NF as official Standards
- 1914: Harrison Narcotic Act
 - Limited the indiscriminant use of addicting drugs by regulating the importation, manufacture, sale, and use of opium, cocaine, marijuana, and their compounds or derivatives

Laws Affecting Drug Administration (Cont.)

- 1938: Federal Food, Drug, and Cosmetic Act
 - Empowered the FDA to pre-market safety standards for drugs. Amended in 1951 by the *Durham-Humphrey Amendments* to require written or verbal prescriptions from a physician to dispense certain drugs

Laws Affecting Drug Administration (Cont.)

- 1970: Comprehensive Drug Abuse Prevention and Control Act
- AKA Controlled Substances Act
 - Classified drugs by abuse potential and medical usefulness.
 - Regulated manufacture, distribution, and sale of controlled substances.
 - Replaced the Harrison Narcotic Act of 1914

Schedules of Drugs

- Schedule I
 - High abuse potential. No accepted medical indications. (Research use only.)
 - Examples: Heroin, LSD
- Schedule II
 - High abuse potential. Accepted medical indications. Written prescriptions required.
 - Examples: Opium, cocaine, morphine, codeine, oxycodone, methadone

Schedules of Drugs (cont.)

- Schedule III
 - Less abuse potential than schedule II or II; may lead to moderate or low physical dependence
 - Limited opioid amount or combined with noncontrolled substances
 - Examples: Vicodin, Tylenol w/ codeine

Schedules of Drugs (cont.)

Schedule IV

- Low abuse potential compared to schedule III. Limited psychosocial and/or physical dependence
- Examples: Diazepam, lorazepam, phenobarbital

Schedule V

- Lower abuse potential than schedule IV. May lead to psychosocial and/or physical dependence
- Limited amounts opioids; often for cough or diarrhea

Drug Profiles

- Components
 - Names
 - Classification
 - Mechanisms of Action
 - Indications
 - Pharmacokinetics
 - Side effects



- Routes of Administration
- Contraindications
- Dosage
- How supplied
- Special considerations

Patient Care with Medications

- Know the Drug profile
- Practice proper technique
- Know how to observe and document effects
- Maintain current knowledge in pharmacology
- Establish and maintain professional relationships with other health care providers

Patient Care with Medications (cont.)

- Understand pharmacokinetics & pharmacodynamics
- · Have current drug references available
- Take careful drug histories
- Evaluate the compliance, dosage, and adverse reactions
- Consult with medical direction when appropriate

Six Rights of Medication Administration

- Right medication
- Right dose
- Right time
- Right route
- Right patient
- Right documentation

- Pediatric Patients
 - Neonates (Infants from birth to 4 weeks)
 metabolism and excretion may be impaired
 - Children up to one year have diminished plasma protein concentrations. Results in higher free drug availability with drugs that bind to proteins



- Many factors cause a pediatrics drug function to differ radically from an adults
- The Broselow tape primarily addresses drugs administered in the critical care setting



- Geriatric Patients
 - Common physiological effects of aging
 - Cardiac output
 - Renal function
 - Brain mass
 - Total body water
 - Body fat
 - Serum albumin
 - Respiratory capacity

- These changes can lead to:
 - Altered pharmacodynamics & pharmacokinetics
 - Decreased rates of metabolism and excretion
 - Decreased protein binding because of decrease level of serum albumin
- Result Dosages may have to be decreased
- Elderly also suffer from multiple disease processes
- May be on chronic medications that can affect emergency medications

- **Pregnant Patients**
 - Anatomical & Physiological changes
 - Increased cardiac output
 - Increased heart rate
 - Increased blood volume (up to 45%)
 - Decreased protein binding
 - Decreased hepatic metabolism
 - Decreased blood pressure



- Drug has the potential to cross the placenta and affect the fetus
- Drug therapy can affect a breast-feeding infant

Paramedics Responsibilities in Administration of Medications

- Paramedics are personally, legally, morally and ethically responsible for the safe administration of medications
 - Know the precautions and contraindications for all medications you administer
 - Practice proper technique
 - Know how to observe and document drug effects

Paramedics Responsibilities in Administration of Medications

- Maintain a current knowledge in Pharmacology
- Establish and maintain professional relationships with other health care providers
- Understand the pharmacokinetics and pharmacodynamics
- Have current medication references available

Paramedics Responsibilities in Administration of Medications

- Take careful drug histories including:
 - Name, strength, and daily dose of prescribed drugs
 - Over-the-counter drugs
 - Vitamins
 - Herbal medications
 - Folk-medicine or folk remedies
 - Allergies
- Evaluate the compliance, dosage, and adverse reactions
- Consult with medical direction when appropriate

Pharmacokinetics

- Strictly defined, pharmacokinetics is the study of the basic processes that determine the duration and intensity of a drug's effect
- Pharmacokinetic Processes
 - Absorption
 - Distribution
 - Biotransformation
 - Elimination

Physiology of Transport

- Active transport
 - Requires the use of energy to move a substance
- Carrier-mediated diffusion
 - AKA Facilitated Diffusion
 - Process in which carrier proteins transport large molecules across the cell membrane

Physiology of Transport (cont.)

- Passive transport
 - Movement of a substance without the use of energy
- Diffusion
 - Movement of solute in a solution from an area of higher concentration to an area of lower concentration

Physiology of Transport (cont.)

Osmosis

Movement of solute in a solution from an area of lower solute concentration to an area of higher solute concentration

Filtration

 Movement of molecules across a membrane from an area of higher pressure to an area of lower pressure.

Absorption The process of movement of a drug from the site of application into the body and into the extra-cellular compartment.

Absorption

- Affected by many factors including:
 - Solubility of the drug
 - Concentration of the drug
 - pH of the drug/pH of the patient
 - Site of absorption
 - Absorbing surface area
 - Blood supply to the site of absorption

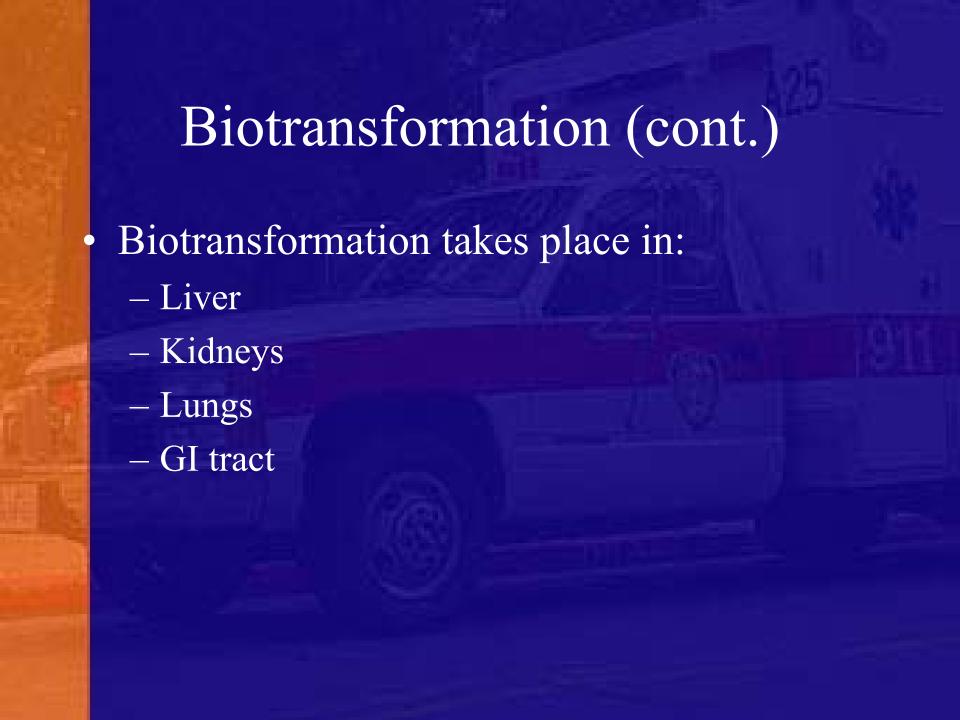
Distribution

- The process whereby a drug is transported from the site of absorption to the site of action
- Affected by several factors:
 - Cardiovascular function
 - Regional blood flow
 - Drug storage reservoirs



Biotransformation

- A special name for metabolism
- The body's breaking down of chemicals
 - Has one of two effects on drugs
 - It can transform the drug into a more or less active metabolite
 - It can make the drug more water soluble (or less lipid soluble) to facilitate elimination



Biotransformation (cont.)

- First-pass effect
 - Blood supply from the GI Tract passes through the liver before moving on through the systemic circulation.
 - First pass may completely inactivate many drugs
 - These drugs must be given IV rather than orally

Biotransformation (cont.)

- Biotransformation begins immediately following introduction of the drug
 - Certain drugs are rapidly transformed
 - Epinephrine is active as administered and rapidly metabolized to inactive forms

Biotransformation (cont.)

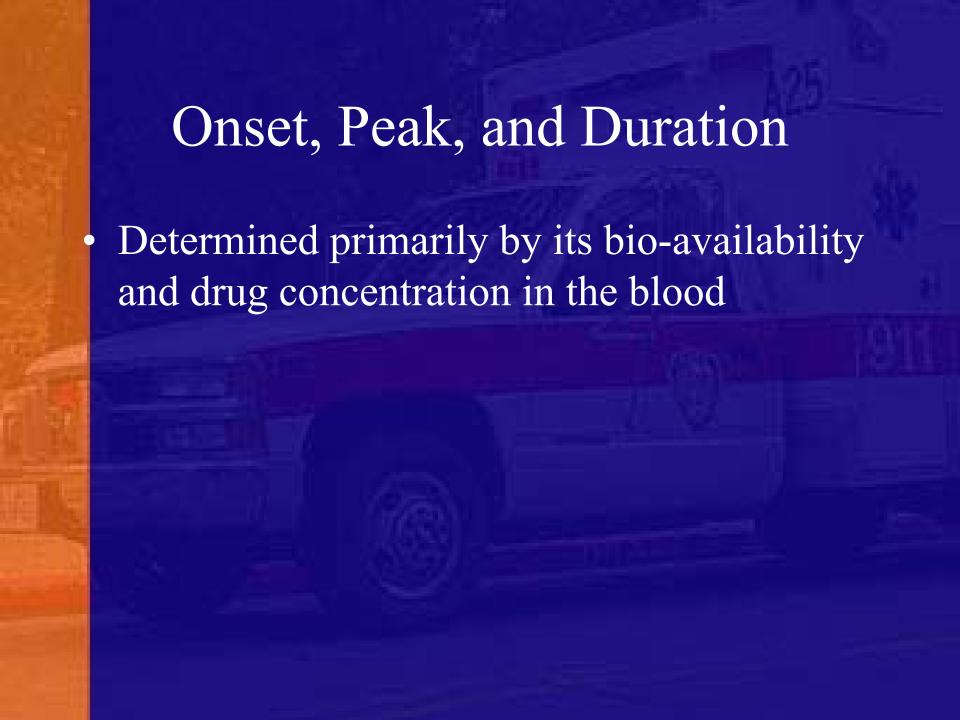
- The liver's microsomal enzymes react with drugs in two ways
 - Phase-I (non-synthetic reactions.)
 - Most often oxidize the parent drug
 - May reduce or hydrolyze the drug
 - Phase II (synthetic reactions.)
 - AKA conjugation reactions, combine the pro-drug or its metabolites with an endogenous chemical, usually making the drug more polar and easier to excrete

Elimination

- Refers to movement of a drug or its metabolites from the tissues back into the circulation and to the organs of excretion
 - Urine Via the Kidneys
 - Bile Via the Liver
 - Feces Via the Intestines
 - Expired air Via the Lungs
 - Sweat, saliva, and breast milk



- Eliminated in original form or as metabolites
- Elimination is affected by:
 - Drug half-life
 - Accumulation
 - Clearance





Enteral

- Absorption through the GI tract
- Enteral Routes
 - Oral (PO)
 - Orogastric / nasogastric tube (OG/NG)
 - Sublingual (SL)
 - Buccal
 - Rectal (PR)

Enternal (cont.)

- Advantages
 - Simple; Safe
 - Generally less expensive
 - Low potential for infection
- Disadvantages
 - Slow rate of onset
 - Cannot be given to unconscious or nauseated patients
 - Absorbed dosage may vary significantly

Parenteral Routes

- Broadly defined, any Route outside of the GI tract
- Parenteral Routes
 - Topical/Intradermal/Subcutaneous
 - Intramuscular/Intravenous/Intraosseous
 - Endotracheal/Sublingual injection/Inhalation
 - Umbilical/Vaginal/Rectal



- Solid/Pills/Powders/Tablets
- Suppositories/Capsules/Liquid
- Solutions/Tinctures/Suspensions
- Emulsions/Spirits/Elixirs
- Syrups

Pharmacodynamics:

Is the study of mechanisms by which specific drug dosages act to produce biochemical or physiological changes in the body

Actions of Drugs

Medications can act in four different ways

- 1. Bind to a receptor site
- 2. Change the physical properties of cells
- 3. Chemically combine with other substances
- 4. Alter a normal metabolic pathway

Actions of Drugs (Cont.)

- Binding To A Receptor Site
 - A receptor is a specialized protein that combines with a drug resulting in a biochemical effect
 - Affinity (Force of attraction between a drug and a receptor)
 - Efficacy (A drugs ability to cause the expected response)

Actions of Drugs (Cont.)

- Second messenger
 - Chemical that participates in complex cascading reactions that eventually cause a drug's desired effect
- Down-regulation
 - Binding of a drug or hormone to a target cell receptor that causes the number of receptors to decrease

Actions of Drugs (Cont.)

- Up-regulation
 - A drug causes the formation of more receptors than normal
- Stimulation of A Receptor Site
 - Chemicals that stimulate fall into two broad categories

Actions of Drugs (cont.)

- Stimulation of A Receptor Site
 - Chemicals that stimulate fall into two broad categories
 - Agonist
 - Causes it to initiate the expected response
 - Antagonist
 - Causes the drug not to initiate the expected response
 - Some drugs do both--Called agonist-antagonist AKA Partial agonist

Stimulation of A Receptor Site

- Competitive antagonism
 - One drug binds to a receptor and causes the expected effect while also blocking another drug from triggering the same receptor
- Non-Competitive antagonism
 - The binding of an antagonist causes a deformity of the binding site that prevents an agonist from fitting and binding



Other Actions of Drugs

- Changing Physical Properties
 - Osmotic balances across membranes are good examples. (ie: Mannitol)
- Chemically combining with other substances
 - Drugs that participate in chemical reactions that change the chemical nature of their substrates



- Altering a normal metabolic pathway
 - The anticipated product will not form, or, if formed, will be substantially or completely inactive

Responses to Drug Administration

- Side effect
- Allergic reaction
- Idiosyncrasy
- Tolerance
- Cumulative effect
- Drug dependence
- Potentiation

Drug Response Relationship

- Correlates different amounts of drug to the resultant clinical response
- Plasma-level profile
 - Describes the lengths of onset, duration, and termination of action, as well as the drug's minimum effective concentration and toxic levels

Drug Response Relationship (cont.)

- Factors Altering Drug Response
 - Age
 - Body Mass
 - Sex
 - Environment
 - Time of Administration
 - Pathologic state
 - Genetic factors
 - Psychological factors

Drug Interactions

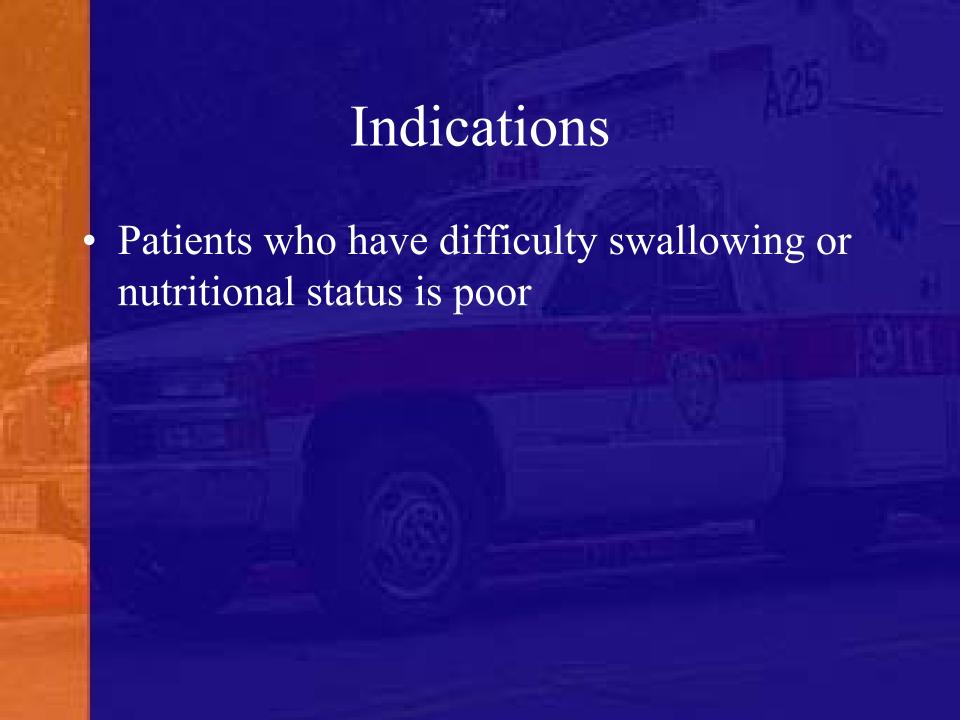
- Variables that may cause drug-drug interactions
- One drug could alter the rate of intestinal absorption.
- The two drugs could compete for plasma protein binding, resulting in one's accumulation at the other's expense

Drug Interactions

- One drug could alter the other's metabolism, thus increasing or decreasing either's bioavailability.
- One drug's action at a receptor site may be antagonistic or synergistic to another's.
- One drug could alter the other's rate of excretion through the kidneys

Drug Interactions One drug could alter the balance of electrolytes necessary for the other drug's expected result







- A 100 ml Cone-tipped syringe
- A 30-50 ml Cone-tipped syringe for medication.
 - 50-100 ml of Normal Saline

Technique

Confirm proper tube placement.



Withdraw the plunger while observing for the presence of gastric fluid or contents.



Instill the medication into the gastric tube.



Gently inject the saline.



Clamp off the distal tube.





Administering medications rectally

Drugs given rectally do not pass through the liver and therefore do not undergo hepatic alteration (first pass effect)

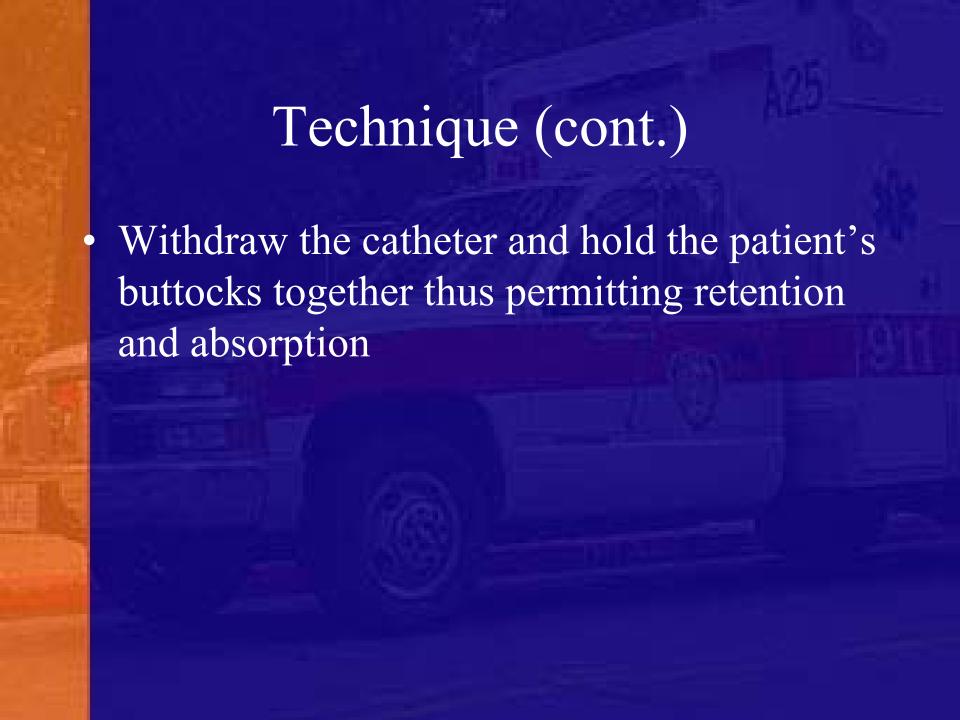
Technique

- Confirm the indication for administration and dose, and draw the correct quantity of medication into a syringe.
- Place the hub of a 14-gauge Teflon catheter (removed from the angiocatheter) on the end of a needleless syringe.

Technique (cont.)

Insert the Teflon catheter into the patient's rectum and inject the medication in the lower part of the rectum.

Administration higher in the rectum may result in the medication's being absorbed by veins that deliver the drug to the portal circulation.



Catheter placement on needleless syringe.



Syringe attached to endotracheal tube.



Prepackaged enema container.







Preparedness

- Have a small sharps container in your jump kit or drug box.
- Make sure all sharps container on ambulance and kit are not filled and lids are in good working condition.
 - Properly close and replace any sharps containers needing to be changed

Preparedness (cont.)

- Sharps disposal must be completed immediately after administrating medication
- Never lay sharps on the ground or stick them into the ground.
- Never stick sharps into the bench seat or any other surface in the ambulance



- Needle handling precautions
- Minimize task in a moving ambulance.
- Properly dispose of all sharps
- Recap needles only as a last resort